Short Report

Phlebology

Management of isolated distal deep venous thrombosis

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Keywords

Deep vein thrombosis, anticoagulation, duplex ultrasound

Introduction

Venous thromboembolism is the third most common cardiovascular condition, with an annual incidence estimated at 1 in 1000 persons. Of these, a third present with deep venous thrombosis (DVT), with symptoms including localised pain, swelling and tenderness of the calf or thigh, and signs including erythema and warmth in the affected limb. DVTs may result in devastating complications; more acutely, DVTs can result in pulmonary embolism which is associated with high morbidity and mortality rate. The longer-term sequelae of DVTs include post-thrombotic syndrome (PTS), with chronic limb swelling and ulceration leading to prolonged quality of life impairments.

Of all lower limb DVTs diagnosed on ultrasound, between 30 and 50% may be classified as an isolated distal DVT (IDDVT).² IDDVTs refer to thrombosis of the infrapopliteal veins, which include the inter-muscular deep veins (including the posterior tibial, anterior tibial, and peroneal veins) and the calf intra-muscular veins (gastrocnemius and soleal veins). These are usually associated with transient risk factors^{2,3} and exhibit lower risk for propagation. However, IDDVTs are still associated with the complications discussed above, with a 17% PTS rate, one in 50 developing ulceration after experiencing an IDDVT,⁴ and around 5% risk of developing a pulmonary embolism.⁵ Appropriate management of IDDVTs is therefore essential to prevent patients from experiencing long-term sequelae stemming from this condition.

In general, IDDVTs may be treated conservatively with compression and/or serial imaging and observation, or with anticoagulation to help limit proximal extension and recurrence. However, there remains significant debate on selecting the optimal treatment for specific patients presenting with IDDVTs. This article seeks to highlight the management recommendations from the International Union of Phlebology (UIP), identifying characteristics that may help with patient stratification for different treatment options. The accompanying one-page summary also

provides a management algorithm that can be followed to help clinicians determine the optimal management plan for patients presenting with IDDVTs (Figure 1).

Management recommendations

It is recommended that patients presenting with IDDVTs should be stratified based on their symptom burden (high or low burden) and risk factors for proximal extension (Table 1).⁶ Two main treatment strategies exist: conservative management with serial imaging and regular follow-up, or anticoagulation together with compression therapy. IDDVT, if presenting without severe symptoms or risk factors for extension, can be managed conservatively. Serial imaging of the deep veins for 2 weeks is recommended, with regular follow-up with a vascular specialist for monitoring. If there are signs of proximal extension or recurrence, the patient will then be started on anticoagulation and compression.

However, should there be significant symptoms such as calf pain or any risk factors for thrombus extension, patients should be further stratified according to their risk of bleeding. If they are deemed to be at low risk of bleeding, 3 months of anticoagulation may be prescribed together with compression therapy. Patients should also be advised

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Management of Isolated Distal Deep Venous Thrombosis (DVT)

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Summary

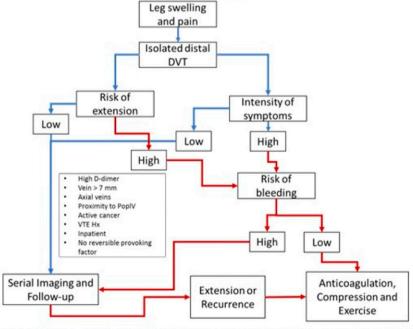
- The need for treatment of isolated distal deep vein thrombosis (DVT) remains controversial.
- Distal DVT includes thrombosis of inter-muscular deep veins (posterior tibial, anterior tibial and peroneal veins) and calf muscular veins (gastrocnemius and soleal veins).
- Distal DVT is more commonly associated with:
 - · Transient risk factors
 - · Lower propagation rate, lower incidence of pulmonary embolism and lower mortality
 - Lower risk for development of post-thrombotic syndrome

Isolated distal DVTs are treated based on symptoms and risk factors

Management Recommendations:

- Isolated distal DVT without severe symptoms or risk factors for extension serial imaging of the deep veins for 2 weeks
- Significant calf pain or risk factors for thrombus extension anticoagulation for 3 months
- Inter-muscular DVT (posterior tibial, peroneal, or anterior tibial)—treat with either anticoagulation or serial imaging depending on symptoms and ongoing risk factors for thrombus extension
- Intramuscular DVT serial imaging and follow-up

Management Algorithm



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Figure 1. One-page guideline.

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Table I. Risk factors for proximal extension of an IDDVT.

High D-dimer Act
Vein diameter >7 mm His
Axial vein involvement Inp
Proximity to popliteal vein No

Active malignancy
History of VTE
Inpatient admission

No reversible provoking factor

to exercise. However, if they are at a high risk of bleeding, it is recommended that clinicians perform serial imaging and follow-up, and only prescribe anticoagulation if there are signs of extension or recurrence on imaging.⁷

Management may also be further stratified according to the location of the DVT either in the inter-muscular deep veins or the intra-muscular veins. It is recommended that intra-muscular IDDVTs can be managed with serial imaging and follow-up alone. Inter-muscular IDDVTs (in the posterior tibial, peroneal, or anterior tibial veins) may be managed with either anticoagulation or serial imaging depending on patients' symptoms and risk factors for thrombus extension as detailed above.

Discussion

This article provides a one-page clinical practice guideline summarising the clinical management of IDDVT. It is part of a series of publications for the UIP One-Page Guidelines which are aimed at ensuring that patients with venous disease receive timely and appropriate care based on current best evidence and expert consensus.

Despite the frequency of IDDVTs encountered in clinical practice, the current evidence remains sparse and there remains controversy regarding whether IDDVTs require treatment at all. This stems from studies showing low risks of proximal extension and high rates of complete resolution without anticoagulant treatment, and low risks of developing valvular insufficiency at late follow-up. However, few studies have directly observed the effects of treatment in confirmed IDDVTs. For example, the more recent CAC-TUS trial showed no significant impact of therapeutic anticoagulation for proximal extension or thromoboembolism rates related to IDDVTs, but increased risk of bleeding. This trial, however, was underpowered due to early trial termination. The lack of trials directly comparing the various treatment modalities represents a gap in the literature that should be addressed urgently.

This one-page guideline has recommended that high-risk or highly symptomatic patients receiving anticoagulation should continue treatment for at least 3 months, which is consistent with the guidelines from the American College of Chest Physicians. ¹⁰ Studies have however used varying protocols ranging from 6 weeks to 3 months, with a meta-analysis suggesting reduction in recurrence rates when comparing patients who received more than 6 weeks of anticoagulant therapy to those who received 6 weeks only. ¹¹

Optimal duration of anticoagulation, in view of the potential bleeding risks (as observed in the CACTUS trial⁷), needs to be clarified further to maximise therapeutic benefits for patients without significantly increasing their risks.

For healthcare providers, this article and summary document should highlight the complexity of management of IDDVTs. All healthcare staff members, including emergency medicine practitioners and vascular specialists, that are likely to encounter a patient with an IDDVT should be aware of the need for individualised risk assessment and targeted management, with appropriate escalation of therapies applied when required. This is in line with national and international guidelines and further reinforced by the management algorithm provided in this one-page guideline. This will facilitate the urgent assessment and interventions indicated in this population of patients.

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Guarantor

Professor Alun Davies on behalf of the UIP.

Contributorship

K.P. and A.H.D. conceptualised the design of the short report and one-page guideline.

F.L., D-I.K., and T.W. contributed to the literature review and formulation of the recommendations.

M.T. contributed to the formatting and layout of the one-page guideline (Figure 1) and wrote the initial draft of the short report. All authors reviewed the short report prior to submission.

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